Amendments to the claims:

1. (Currently Amended) A composition comprising: extract of a plant *Prunella Linn* ~~or *Rabdosis (Blume) Hasskarl*~~ containing corosolic acid at a concentration of at least 0.01% by weight.

2. (Withdrawn) The composition of claim 1, wherein the concentration of corosolic acid is at least 0.1% by weight.

3. (Original) The composition of claim 1, wherein the concentration of corosolic acid is at least 1% by weight.

4. (Withdrawn) The composition of claim 1, wherein the concentration of corosolic acid is at least 10% by weight.

5. (Withdrawn) The composition of claim 1, wherein the extract is an extract of the whole plant of *Prunella Linn* or *Rabdosis (Blume) Hasskarl*.

6. (Original) The composition of claim 1, wherein the extract is an extract of the portion of the plant that grows above the ground.

7. (Original) The composition of claim 1, further comprising:
ursolic acid, 2α, 19α-dihydricursolic acid or daucosterol.

8. (Original) The composition of claim 1, wherein the corosolic acid is in a form of solid.

9. (Original) The composition of claim 1, wherein the extract is in a form of liquid.

10. (Original) A Pharmaceutically acceptable composition, comprising:
a pharmaceutically acceptable excipient; and
extract of a plant *Prunella Linn* ~~or *Rabdosis (Blume) Hasskarl*~~ containing corosolic acid at a concentration of at least 0.01% by weight.

11. (Original) The composition of claim 10, wherein the pharmaceutically acceptable composition is suitable for oral administration to a human.

12. (Original) The composition of claim 10, wherein the pharmaceutically acceptable composition is formulated with the excipient in a form selected from the group consisting of tablets, pills, dragees, capsules, emulsions, lipophilic and hydrophilic suspensions, liquids, gels, syrups, slurries, and suspensions.

13. (Original) The composition of claim 12, wherein the pharmaceutically acceptable composition is formulated in hard or soft-gel capsules.

14. (Original) The composition of claim 10, wherein the excipient is selected from the group consisting of glycerol, sorbitol, lactose, magnesium stearate, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and polyvinylpyrrolidone.

15. (Original) The composition of claim 10, wherein the excipient is an pharmaceutically acceptable oil.

16. (Original) The composition of claim 15, wherein the pharmaceutically acceptable oil is selected from the group consisting of corn oil, wheat germ oil, soy bean oil, rice bran oil, rapeseed oil, sesame oil, and fish oil.

17. (Original) The composition of claim 10, wherein the concentration of corosolic acid is at least 1% by weight.

18. (Original) The composition of claim 10, wherein the concentration of corosolic acid is at least 10% by weight.

19. (Original) The composition of claim 10, further comprising:

ursolic acid, 2α, 19α-dihydricursolic acid or daucosterol.

20-64. (Canceled without Prejudice)